Supplementary Material

Müller Cells Stabilize Microvasculature through Hypoxic Preconditioning

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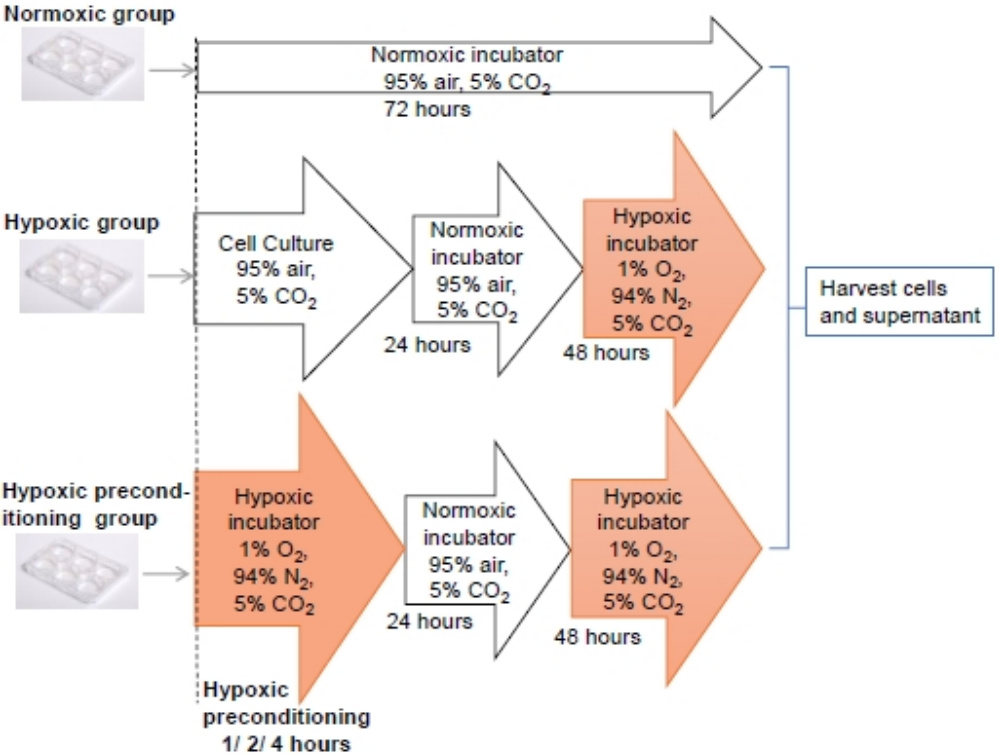
Supplementary Figure 1. Hypoxic preconditioning method. To generate the hypoxic preconditioning condition, the cultures were gassed with 1% O_2, 94.5% N_2, and 5% CO_2, and control cultures were incubated under normoxic conditions for the same duration. After the indicated hypoxic period (1/ 2/ 4 hours), reoxygenation was performed by transferring the cells into a regular normoxic incubator (95% air, 5% CO_2), and cells were incubated for another 24 hours for hypoxia assays



Supplementary Figure 1